

professional meetings or publication in professional journals.
Respondents: Participants in ACF programs being evaluated; participants

in ACF demonstrations; recipients of ACF Grants and individuals served by ACF Grantees; comparison group

members; and other relevant populations, such as individuals at risk of needing ACF services.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Survey development field tests, respondent debriefing questionnaires, cognitive interviews, split sample experiments, focus groups	3,825	1	1	3,825

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Planning, Research, and Evaluation, 330 C Street SW., Washington, DC 20201, Attn: OPRE Reports Clearance Officer. Email address: OPREinfocollection@acf.hhs.gov. All requests should be identified by the title of the information collection.

The Department specifically requests comments on (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Mary Jones,
 ACF/OPRE Certifying Officer.
 [FR Doc. 2017-22718 Filed 10-19-17; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Adoption and Foster Care Analysis and Reporting System (AFCARS).

OMB No.: 0970-0422.

Description: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, the Administration for Children and Families (ACF) is publishing this notice that summarizes the following proposed collection of information for public comment:

Title of Collection: Adoption and Foster Care Analysis and Reporting System (AFCARS).

OMB Control Number: 0970-0422, expiration date February 28, 2018.

Type of Review: An extension of an existing information collection.

Respondents/Affected Public: Title IV-E State, Local, and Tribal Child Welfare Agencies.

There have been no revisions to the existing AFCARS data collection found under federal regulations at 45 CFR 1355.40. The requirements for AFCARS being addressed by this notice are scheduled to end on September 30, 2019. ACF is requesting that OMB approve a three-year renewal to cover the period when States and IV-E Tribes must continue submitting the existing requirements.

On December 14, 2016, a new set of requirements (45 CFR 1355.41-44) for AFCARS reporting was published as a final rule and will go into effect on October 1, 2019. At that time, the requirements covered by this notice (45 CFR 1355.40) will be replaced by the new regulatory requirements. The requirements for the updated AFCARS requirements found in federal regulations at 45 CFR 1355.41-44 are covered by a different OMB control number (0970-0457) and are not addressed by this Notice.

SUPPLEMENTARY INFORMATION: In addition to this Notice, Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or update of an existing collection of information, before submitting the collection to OMB for approval. In compliance with this requirement, ACF published a notice in the **Federal Register** on June 30, 2017 and invited comment on: (1) Whether the proposed collection of information is necessary for the proper performance of ACF's functions, including whether the information will have practical utility; (2) the accuracy of ACF's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques when appropriate, and other forms of information technology for the existing AFCARS reporting that is required by States and Tribes.

ACF received 71 comment letters to the first notice. No substantive comments were received regarding the existing AFCARS data collection in 45 CFR 1355.40 under OMB Number 0970-0457. Instead, commenters provided feedback on the new AFCARS data

collection under 45 CFR 1355.41–.47 to be implemented October 1, 2019. Commenters addressed support for the collection of information under the Indian Child Welfare Act and other areas covered by the new requirements. ACF received no comment on the specific burden hours for the existing AFCARS requirements undergoing renewal.

AFCARS is mandated by 42 U.S.C. 679. The regulation at 45 CFR 1355.40 and the appendices to 45 CFR 1355 set

forth the requirements of section 479 of the Social Security Act for the collection of uniform, reliable information on children who are under the responsibility of the State or Tribal title IV–B/IV–E agency for placement, care, and adoption. The AFCARS requirements under 45 CFR 1355.40 have been in effect since October 1, 1993 for States. In 2009, section 479B(b) of the Act was enacted authorizing direct Federal funding of Indian Tribes, Tribal organizations, and Tribal

consortia that choose to operate a foster care, adoption assistance and, at Tribal option, a kinship guardianship assistance program under title IV–E of the Act. The data collected informs State/Tribal/Federal policy decisions, program management, and responses to Congressional and Departmental inquiries.

Respondents: Title IV–E State and Tribal Child Welfare Agencies.

ESTIMATED TOTAL ANNUAL BURDEN HOURS

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
AFCARS	59	2	2,188	258,215
Estimated Total Annual Burden Hours	258,215

Additional Information: Copies of the regulation containing the data elements may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201. Attention Reports Clearance Officer. All requests should be identified by the title of the information collection. Email address: infocollection@acf.hhs.gov.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Email: OIRA_SUBMISSION@OMB.EOP.GOV, Attn: Desk Officer for the Administration for Children and Families.

Mary Jones,

ACF/OPRE Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2016–D–1229]

Current Good Manufacturing Practice Requirements for Food for Animals; Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a guidance for industry #235 entitled “Current Good Manufacturing Practice Requirements for Food for Animals.” This guidance helps domestic and foreign facilities that are required to register as food facilities under the Federal Food, Drug, and Cosmetic Act (the FD&C Act) determine whether and how they need to comply with the current good manufacturing practice requirements of the Current Good Manufacturing Practice, Hazard Analysis, and Risk-Based Preventive Controls for Food for Animals final rule.

DATES: The announcement of the guidance is published in the **Federal Register** on October 20, 2017.

ADDRESSES: You may submit either electronic or written comments on Agency guidances at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <https://www.regulations.gov>. Follow the instructions for submitting comments.

Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else’s Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <https://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see “Written/Paper Submissions” and “Instructions”).

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in “Instructions.”

Instructions: All submissions received must include the Docket No. FDA–2016–D–1229 for “Current Good Manufacturing Practice Requirements